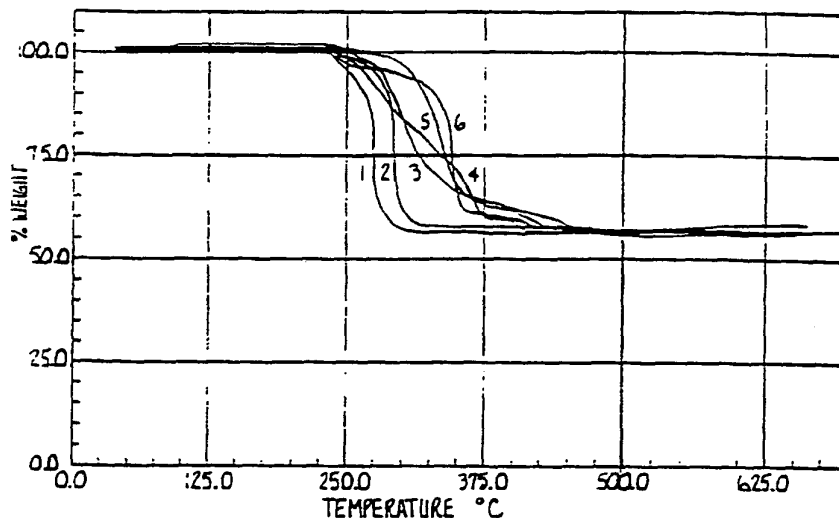




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : C01B 11/14, 11/18	A1	(11) International Publication Number: WO 93/17961 (43) International Publication Date: 16 September 1993 (16.09.93)
(21) International Application Number: PCT/US93/01562 (22) International Filing Date: 23 February 1993 (23.02.93) (30) Priority data: 07/851,171 13 March 1992 (13.03.92) US (71) Applicant: PURITAN-BENNETT CORPORATION [US/US]; 10800 Pflumm Road, Lenexa, KS 66215 (US). (72) Inventors: ZHANG, Yunchang ; 13229 West 107th Terrace, Lenexa, KS 66210 (US). CANNON, James, C. ; 15110 West 123rd Street, Olathe, KS 66062 (US). (74) Agents: COLLINS, John, M. et al.; Hovey, Williams, Timmons & Collins, 1101 Walnut Street, Suite 1400, Kansas City, MO 64106 (US).		(81) Designated States: AU, CA, JP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>

(54) Title: NON-TOXIC CHEMICAL OXYGEN GENERATING COMPOSITIONS

1) 99.7% NaClO₃ + 1% CoO_x
 3) 98.7% NaClO₃ + 1% Li₂B₄O₇ + 1% CoO_x
 5) 97.7% NaClO₃ + 2% Na₂SiO₃ + 1% CoO_x

2) 95.7% NaClO₃ + 4% SrCO₃ + 1% CoO_x
 4) 97% NaClO₃ + 2% Ca₂P₂O₇ + 1% CoO_x
 6) 98% NaClO₃ + 1% Li₃PO₄ + 1% CoO_x

(57) Abstract

An oxygen-generating candle composition is disclosed of the type that includes an amount of an alkali metal chlorate or perchlorate oxygen source and which upon initiation and decomposition yields oxygen and residual chlorine, wherein the improvement comprises a non-toxic additive in the candle composition for suppression of the residual chlorine and to enhance uniform oxygen generation and evolution. The additive is selected from the group consisting of from about 0.05% to about 10% by weight of lithium aluminate, metaborate, tetraborate, phosphate or pyrophosphate, metasilicate or orthosilicate, or carbonate, calcium phosphate or pyrophosphate, strontium carbonate or phosphate, or sodium metasilicate or orthosilicate. A metal oxide catalyst, a metal powder fuel, and a binder or filler are also included in the composition.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Monaco	TC	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam
FI	Finland				

NON-TOXIC CHEMICAL OXYGEN GENERATING COMPOSITIONSBackground of the Invention

5 This invention is broadly concerned with an improved candle for generating breathable oxygen by initiation and decomposition of an alkaline metal chlorate or perchlorate source. During decomposition of the chlorate or perchlorate, toxic free chlorine gas may be evolved. Furthermore, irregular decomposition of the
10 alkali metal chlorate or perchlorate will result in uneven evolution of oxygen.

 Thus, this invention specifically relates to an oxygen generating alkali metal chlorate or perchlorate candle composition having a non-toxic additive compound
15 incorporated therein which regulates and smooths oxygen decomposition and suppresses formation of toxic free chlorine gas.

20 Description of the Prior Art

 Chemical oxygen generating candles containing alkali metal chlorates or perchlorates have long been used as an emergency supply of breathable oxygen where weight and space limitations are important factors. Exemplary in
25 this respect are aircraft which fly at high altitudes, submarines, and other closed atmospheric enclosures which house personnel or passengers.

 Many previous oxygen candles which employed an alkali metal chlorate or perchlorate compound as the oxygen source have utilized barium peroxide (BaO_2) as an
30 additive to smooth out the decomposition rate of the chlorate or perchlorate and to suppress formation of free chlorine which is a toxic gas. A metal oxide catalyst has also commonly been used to facilitate decomposition of the
35 oxygen source, and a powdered metal fuel such as iron

powder incorporated to furnish extra heat to help sustain the reaction process. A small portion of glass powder and/or glass fibers have also been employed as a binder. Although BaO_2 performs the function of suppressing free chlorine formation and smoothing oxygen evolution during decomposition of alkali metal chlorate or perchlorate oxygen sources, it is in and of itself a poisonous compound. Unless extraordinary precautions are taken, tiny BaO_2 particles may be released along with the oxygen generated, presenting an immediate health hazard to the person breathing the oxygen being generated and possibly contaminating the surrounding atmosphere as well. Equally as importantly, environmental regulations require the use of costly procedures and equipment to dispose of scraps and expended oxygen candles containing BaO_2 and/or its reaction products, such as BaCl_2 .

Several alkali metal oxides such as Na_2O , Na_2O_2 and KO_2 have also been suggested as catalysts for NaClO_3 decomposition and to suppress acidic chlorine gas formation, but have not met with widespread usage.

M.M. Markowitz et al., in U.S. Patent No. 3,293,187 and in an article in I&EC Product Research and Development, 3(4), 323 (1964) disclosed that Li_2O_2 and other alkaline compounds can suppress free chlorine formation in lithium perchlorate based oxygen candles. Manganese powder was used as a fuel, and manganese oxides formed during manganese oxidation were suggested as catalysts. In view of the fact that decomposition of such lithium perchlorate compositions tends to produce relatively high concentration of chlorine gas, Markowitz had to use 5% Li_2O_2 by weight. With high lithium peroxide loading the integrity of the candle remains a problem since the candle tends to melt and flow away.

U.S. Patent Nos. 3,702,305 and 3,736,104 disclose use of sodium oxide and peroxide catalyst in a

sodium chlorate oxygen candle. However, these alkali metal oxides are air sensitive and caustic. They will absorb moisture and CO_2 from the air and decompose, which results in decomposition upon exposure to air with concomitant production, use and storage problems. Candles using such compounds must be produced by a dry process, and production by wet processing is precluded.

Chinese Patent No. 1035248A describes the use of lithium peroxide as a chlorine suppressant in a sodium chlorate oxygen candle. The formulation contains 2%-7% cobalt oxide, 0%-5% MnO_2 , 1%-6.5% Li_2O_2 and 1%-8% kaolin. The ingredients were dry mixed and pressed. The total oxygen generation is 6 liters. Since magnesium is a very active metal and it can react with water and release hydrogen, the procedure is not practical if carried out as a wet process. Magnesium is also relatively difficult to ignite. Furthermore, because a relatively high proportion of lithium peroxide is used, a similar high proportion of cobalt oxide must be employed and several percent of kaolin has to be incorporated to maintain the integrity of the candle. High catalyst and additive loading increases candle weight and lowers the oxygen yield.

Cobalt oxide and lithium peroxide are also relatively expensive chemicals. High loading significantly increases the cost of the candle. The Chinese patent also describes low fuel or no fuel candle compositions. However, no fuel or very low fuel candles are generally more sensitive to environmental temperatures, and have difficulty functioning in cold environments.

Summary of the Invention

The present invention overcomes the problems noted above and provides a greatly improved oxygen-generating composition which contains no poisonous compounds such as barium peroxide or other toxic compounds, or highly unstable and caustic compounds such as sodium oxide and peroxide and potassium superoxide. The new formulations are suitable for either wet or dry processing. The oxygen candle additives described herein are non-toxic and reasonably stable in air and function both to smooth the sodium chlorate decomposition rate, and to suppress chlorine formation. The additive enhanced alkali metal chlorate or perchlorate formulations function well across a wide range of environmental temperatures.

Broadly speaking, the oxygen generating candle includes an oxygen source which upon initiation and decomposition, yields oxygen, with a non-toxic additive being incorporated into the candle to smooth oxygen evolution and to suppress formation of residual chlorine gas resulting from the side reactions taking place. The additive of the present invention includes one or more alkaline compounds selected from the group consisting of (a) a compound of the formula $M_aH_bX_cO_d$ where M = an alkali metal or alkaline earth metal, a = a number which is one or more, H = hydrogen, b = a number which is 0, or one or more, X = an element chosen from the group consisting of the elements of Group IIIA, the elements of Group IVA, the elements identified by the numbers 15 and 51 of Group VA, the elements of Group IVB, the elements of Group VB, the elements of Group VIB, and the element identified by the number 25 of Group VIIB of the Periodic Table, c = a number which is one or more, O = oxygen, and d = a number which is one or more; (b) lithium, sodium, potassium, rubidium, cesium, magnesium and strontium hydroxides; (c) magnesium, calcium and strontium peroxides; and (d)

mixtures of the foregoing.

In particularly preferred forms the candle includes: an alkali metal chlorate or perchlorate oxygen source; from about 0.05% to about 10% by weight of lithium aluminate, metaborate, tetraborate, phosphate or pyrophosphate, metasilicate or orthosilicate, or carbonate, calcium phosphate or pyrophosphate, strontium carbonate or phosphate, or sodium metasilicate or orthosilicate; a metal oxide catalyst; a metal powder fuel; and a binder or filler.

When used in aircraft, weight is one of the most important factors to be considered. Therefore, Li_2O_2 , LiAlO_2 , LiBO_2 , $\text{Li}_2\text{B}_4\text{O}_7$, Li_2SiO_3 , Li_3PO_4 and Li_2SiO_3 are preferred because of their relatively lower molar weight. Substitution of an equimolar amount of one of these lithium compounds for BaO_2 reduces the overall candle weight.

It is also an objective to make an oxygen candle which can operate over a wide range of surrounding temperatures. Oxygen candles used in aircraft must function well in a wide temperature band which may range from -30°C to 60°C . If the candle burns substantially faster at higher surrounding temperatures, a much larger candle must be used in order for the candle to supply oxygen for a required duration over the requested wide range of temperatures.

A sufficient amount of the alkaline additive is introduced into the candle body composition to increase the decomposition temperature of the alkali metal chlorate or perchlorate oxygen source to a level above the melting temperature thereof that assures steady oxygen generation.

Brief Description of the Drawings

Fig. 1 depicts temperature programmed decomposition profiles of sodium chlorate catalyzed with cobalt oxide, and sodium chlorate compositions catalyzed with cobalt oxide and decomposition smoothing and chlorine inhibiting agents comprising strontium carbonate, lithium tetraborate, calcium pyrophosphate, sodium silicate and lithium phosphate respectively; and

Fig. 1A shows similar decomposition curves for sodium chlorate alone, and sodium chlorate catalyzed with cobalt oxide and containing calcium hydroxide, lithium carbonate, lithium peroxide, barium oxide and lithium metaborate respectively;

Fig. 2 depicts the effect of cobalt oxide on sodium chlorate decomposition, and illustrates the decomposition curves for sodium chlorate catalyzed with cobalt oxide, and sodium chlorate catalyzed with cobalt oxide and containing sodium bicarbonate, sodium carbonate, sodium chromate, potassium hydroxide, and sodium phosphate respectively; and

Fig. 2A is a similar depiction of the decomposition curves for sodium chlorate catalyzed with cobalt oxide and containing sodium hydroxide, barium hydroxide, lithium hydroxide, sodium metaborate and strontium hydroxide respectively;

Fig. 3 is a graphical representation of the rate of flow of oxygen from a candle composition as set forth in Example 1;

Fig. 4 is a graphical representation of the rate of flow of oxygen from a candle composition as set forth in Example 2;

Fig. 5 is a graphical representation of the rate of flow of oxygen from a candle composition as set forth in Example 3;

Fig. 6 is a graphical representation of the rate

of flow of oxygen from a candle composition as set forth in Example 5;

Fig. 7 is a graphical representation of the rate of flow of oxygen from a candle composition as set forth in Example 6; and

Fig. 8 is a cross-sectional view of a chemical oxygen generator useful in producing oxygen utilizing an alkali metal chlorate or perchlorate candle composition in accordance with the present invention, with a typical candle block being illustrated in cross-section in the generator canister.

Description of the Preferred Embodiment

Referring now to the drawings, a representative chemical oxygen generator 10 that may be used in accordance with the present invention is illustrated in Fig. 8 and includes candle 12 installed in a canister 14 along with suitable ignition means 16. Those skilled in the art will appreciate that the candle of the invention may be used in connection with any number of other oxygen generation devices.

Canister 14 includes a tubular housing 18 closed by end caps 18a and 18b, an interior core locator partition 20, spatter shield 22, outlet valve 24 and a pair of relief valves 26, 28 mounted on end cap 18a. It is to be understood that the number and location of outlet and relief valves may vary with the overall size and intended use of the apparatus. Conventional filter material (not shown) occupies the space between core locator 20 and end cap 18a. Candle 12 is surrounded by thermal insulation 32, 34. Mounting lugs 36, 38, 40 are provided on the exterior of the canister surface.

Ignition means 16 includes pull pin 42, spring 44, hammer 46 and primer 48. Although a percussion type

ignition means is preferred, any suitable form of ignition means may be employed.

Candle 12 broadly includes body 50 and an ignition pellet 52. In order to match various high altitude descent oxygen flow specifications, the preferred candle 12 has a weight of between 222 grams and 228 grams and is 5.0 inches long with a diameter of 1.4 inches at the ignition end and 1.1 inches at the outlet end. The preferred ignition pellet 52 has a diameter of 0.56 inches, is 0.25 inch thick and weighs 1.5 grams. Body 50 consists of an oxygen source and a nontoxic additive compound incorporated into the candle. Candle 12 may be formulated to further include a catalyst, a metal powder fuel, and a binder or filler. The candle 12 may be constructed of multiple layers of varying chemical composition. As shown in Fig. 8, the candle may include a first composition layer 54 in contact with ignition pellet 52, and second composition layer 56 and layers thereafter including a nontoxic additive as described herein. In other embodiments, the first composition layer 54 may also contain the nontoxic additive. The preferred candle includes additional layers 58, 60, 62, although any number of layers may be used.

In operation, a user engages pull pin 42 either directly or by means of a cord or other similar remote activation mechanism. When pin 42 is pulled, spring 44 is actuated, causing hammer 46 to hit primer 48 and effect ignition. Primer 48 ignites pellet 52 loaded at the top of the candle body 50. The heat generated from ignition pellet 52 then initiates reaction and therefore decomposition of the candle body. Once initiated, the candle will sustain an exothermic decomposition reaction at a uniform rate to release oxygen at a stable flow rate until the chemical core has completely reacted.

The gaseous reaction products pass through

filter material 30 to remove solid particles and any residual chlorine and carbon monoxide and thence through outlet valve 24 which may be connected to a gas manifold or other oxygen delivery means. In the event of blockage of outlet valve 24 or other increased pressure inside canister, relief valves 26, 28 allow gases to escape. The generated oxygen which passes through outlet valve 24 has the high purity required for breathing. In preferred embodiments the chlorine level is lower than 50 ppb, carbon monoxide level is lower than 3 ppm and carbon dioxide is lower than 1500 ppm. The candle operates well over wide environmental temperatures.

Alkali metal chlorates or perchlorates such as NaClO_3 , KClO_3 , LiClO_3 , KClO_4 , LiClO_4 and NaClO_4 or mixtures thereof can be used as oxygen generating sources. The preferred chlorate is sodium chlorate and the preferred perchlorate is potassium perchlorate.

Sodium chlorate is a particularly preferred oxygen source because it has a relatively high oxygen yield per unit weight compared to potassium chlorate and a reasonably low decomposition temperature compared to potassium chlorate and perchlorate and lithium perchlorate. Decomposition of sodium chlorate is exothermic once initiated, which is essential for self-sustaining operation of the candle. The price of sodium chlorate is also lower than other alkali metal chlorates and perchlorates.

One or more alkaline compounds of lithium, sodium, potassium, rubidium, cesium, magnesium, calcium, and strontium may be employed as additives to smooth out the chlorate decomposition rate and to suppress free chlorine formation. Lithium aluminate, lithium carbonate, lithium borates, lithium phosphates, lithium silicates, calcium phosphates and pyrophosphates, strontium carbonate, and strontium phosphate are particularly

preferred additives because they are stable and have low solubility in water and do not readily hydrate. Sodium aluminate, sodium borates and sodium silicates are also preferred because of their stability and low price.

5 Temperature programmed decompositions were conducted using a thermal balance such as the Model 409 Thermal Analyzer available from NETZSCH of Germany. The ingredients for each test were intimately mixed by grinding. Approximately 100 mg of the mixture was then
10 heated to raise the temperature thereof at a rate of 20°C/min in an oxygen stream. The weight of each sample at an elevated temperature was compared against its original weight at room temperature. The weight of each sample was recorded and presented as a percentage of its
15 original weight against its temperature presented in Figures 1, 1A, 2 and 2A, representing the results of the thermal gravimetric analyses using the Model 409 Analyzer.

 In Figure 1, the temperature programmed decomposition profile of NaClO_3 mixed with 1.0% by weight of cobalt oxide catalyst is represented by the line identified by the numeral 1. Cobalt oxide catalyst is represented as CoO_x where x = a number from 1.0 to 1.5. Line 2 is the graphical profile representation for decomposition of sodium chlorate containing 1.0% cobalt
20 oxide and 4.0% strontium carbonate. Line 3 is the graphical representation for sodium chlorate decomposition containing 1.0% lithium tetraborate and 1.0% CoO_x . Line 4 depicts the decomposition curve for sodium chlorate catalyzed with 1.0% CoO_x and containing 2.0% calcium
25 pyrophosphate. Line 5 indicates the decomposition of sodium chlorate catalyzed with 1.0% CoO_x and containing 2.0% sodium silicate. Line 6 shows the results of sodium chlorate decomposition catalyzed with 1.0% CoO_x and 1.0% lithium phosphate.

35 Fig. 1A graphically depicts the decomposition

curves for sodium chlorate alone as indicated by the line 12 in that figure, while lines 7 to 11 inclusive show decomposition curves for sodium chlorate catalyzed with 1.0% CoO_x and also containing as additives 1.0% calcium hydroxide (line 7), 1.0% lithium carbonate (line 8), 0.5% lithium peroxide (line 9), 4.0% barium oxide (line 10) and 1.0% lithium metaborate (line 11) respectively.

Line 1 and line 12 in Figs. 2 and 2A depict the decomposition profiles of NaClO_3 catalyzed by 1.0% CoO_x and NaClO_3 alone respectively. Lines 2-11 in Figs. 2 and 2A depict the decompositions curves of NaClO_3 catalyzed by 1.0% CoO_x and containing 2% NaHCO_3 , 1% Na_2CO_3 , 2% $\text{Na}_2\text{Cr}_2\text{O}_7 \cdot 2\text{H}_2\text{O}$, 3% KOH , 3% $\text{Na}_3\text{PO}_4 \cdot 12\text{H}_2\text{O}$, 3% NaOH , 4% $\text{Ba}(\text{OH})_2$, 2% LiOH , 2% NaBO_2 , and 2% $\text{Sr}(\text{OH})_2 \cdot 8\text{H}_2\text{O}$ respectively.

The weight losses of the samples represented by curves 2-8 and 11 of Figs. 1, 1A and curves 2-11 of Figs. 2, 2A occurred at temperatures higher than the melting temperature of NaClO_3 , which indicates that all these alkaline additives are inhibitors for catalyzed NaClO_3 decomposition. The alkaline additives referred to in Figs. 1 and 1A have all been proven to smooth oxygen evolution and suppress chlorine formation for a sodium chlorate candle. It can be seen that the curves presented in Figs. 2 and 2A are very close to those presented in Figs. 1 and 1A. Therefore, it can be recognized that the alkaline additives referred to in Figs. 2 and 2A will smooth oxygen evolution and suppress chlorine formation in a similar manner.

As is evident from curve 12 as illustrated in Figure 1A, decomposition of the sodium chlorate started at 480°C and was complete at 590°C . When 1.0% cobalt oxide was mixed, the decomposition started at 225°C and completed at 300°C as depicted in curve 1 of Fig. 1 compared to the melting temperature 260°C for NaClO_3 . A considerable portion of the NaClO_3 decomposed in solid

state.

Nonuniform dispersion and irregular contact of the catalyst and NaClO_3 can cause a irregular oxygen evolution. When in solid phase, the oxygen produced has to build up pressure and force its way out from inside of a candle core, which can also result in irregular oxygen flow. In addition, the pressure build up can rapture the candle core and result in cracks and thus produce more erratic operation for the candle.

If NaClO_3 is made to decompose above its melting temperature 260°C , there is much better contact between the catalyst and liquid NaClO_3 . Nonuniform density distribution resulting from uneven pressure during the pressing is eliminated. Irregular oxygen evolution resulting from minor cracks is also unimportant since the liquid NaClO_3 can fill the cracks. Liquid sodium chlorate has a low viscosity. Oxygen produced inside the liquid phase can get out readily. All these effects contribute to a smoother oxygen evolution.

BaO_2 and relatively high levels of Li_2O_2 have been used in NaClO_3 candles to suppress chlorine formation as previously indicated. Curves 9 and 10 in Figure 1A are the decomposition profiles of NaClO_3 with 1.0% cobalt oxide and 0.5% Li_2O_2 and 4.0% BaO_2 respectively. Although the lithium peroxide and barium oxide brought the decomposition above NaClO_3 's melting temperature and thus contributed to smooth oxygen evolution, the disadvantage of BaO_2 as an additive has been described, and the adverse consequences of high Li_2O_2 levels has also been explained.

The additives referred to in Figs. 2 and 2A are less preferred than those set forth in Figs. 1 and 1A because the former are less stable, exhibit some hygroscopicity, or contain water. Furthermore, the additives of Figs. 2 and 2A are somewhat harder to disperse than those referred to in Figs. 1 and 1A.

It is clear from Figs. 1, 1A, 2 and 2A that alkaline compounds of lithium, sodium, potassium, calcium, strontium, and barium all have an inhibiting effect on catalyzed NaClO_3 decomposition and thus can smooth the oxygen evolution and suppress chlorine formation. Rubidium and cesium are very close to sodium and potassium and magnesium is very close to calcium and strontium chemically. Therefore, basic compounds of rubidium, cesium and magnesium can work equally as well.

The inhibiting effect depends on the alkalinity of the additives. Higher alkalinity additive usually provides more inhibiting effect. In basic media chlorine tends to disproportionate into chlorate and chloride especially at elevated temperatures. Therefore, these basic additives can suppress chlorine by either directly reacting with any chlorine formed or by making the candle basic and keeping chlorine from being formed, as a result of shifting of the equilibria involved in chlorate disproportionation.

Additives with small particle size or high surface area are preferred because smaller particle can provide more intimate mixing and thus provide more inhibiting effect. Soluble alkaline compounds such as sodium silicate and borates have more effect because once dissolved the solution can interact with cobalt oxide and sodium chlorate more readily. But once dissolved, these compounds are somewhat more difficult to disperse and a certain degree of hydration and hydrolysis can occur, and thus makes it much harder to dry the candle.

As is evident from Figs. 1, 1A, 2 and 2A, borates, carbonates, chromates, hydroxides, peroxides, phosphates and silicates of the defined alkali and alkaline earth elements are functional to smooth oxygen evolution and suppress chlorine. Therefore, it is manifest that other alkaline compounds of the said

elements such as aluminates, manganates, stannates, titanates, tungstates and zirconates will function in the same manner. Basic salts containing complex anions such as polyphosphates and polysilicates are also useful as alkali metal chlorate or perchlorate candle additives for the same purposes as described previously.

By analogy to sodium bicarbonate (NaHCO_3), other acid salts of the elements described previously can also be used to smooth the decomposition and suppress chlorine. For example, acid salts such as M_2HPO_4 ($\text{M} = \text{Li}, \text{Na}, \text{K}, \text{Rb}, \text{Cs}$) and NHPO_4 ($\text{N} = \text{Mg}, \text{Ca}, \text{Sr}$) are basic and can thus be used. Monobasic phosphates MH_2PO_4 and $\text{N}(\text{H}_2\text{PO}_4)_2$ are generally acidic. But these acidic monobasic phosphates can decompose at relatively low temperature to form a basic salt and thus can also be used. For example, $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$ loses 2 H_2O at 100°C , and decomposes to form $\text{Na}_2\text{H}_2\text{P}_2\text{O}_7$ at 200°C , and form $\text{Na}_3\text{P}_3\text{O}_9$ between 400°C and 500°C . These acid salts are generally less preferred because they tend to produce water upon decomposition.

The decomposition temperature of sodium chlorate decreases with increasing cobalt oxide loading and increases with increasing alkaline compound loading. By using a proper combination of cobalt and the alkaline compounds, it is possible to make sodium chlorate decompose at any given temperature between approximately 225°C and 500°C .

The inhibiting effect of these described alkaline additive compounds are less important to candles which employ alkali chlorates or perchlorates other than sodium chlorate as the oxygen source. For example, the melting temperature of LiClO_3 and LiClO_4 are 129°C and 247°C respectively. The lithium compounds always decompose in liquid phase even when catalyzed by cobalt oxide with no alkaline additive added. An inhibitor is therefore not essential. The melting points of KClO_3 ,

KClO₄ and NaClO₄ are 357°C, 580°C and 468°C respectively. Making them decompose above their melting temperature requires a much higher fuel loading to sustain the decomposition and results in hotter generator. Again,
5 alkaline inhibitors are not as essential when any of these are used alone.

In the case of NaClO₃ mixed with the other alkali metal chlorates or perchlorates, however, the alkaline compounds work in the same way as for NaClO₃ by itself.
10 Examples are NaClO₃/LiClO₄ and NaClO₃/KClO₄ mixtures. NaClO₃ can form eutectic with LiClO₄ or KClO₄. The melting temperatures of the eutectics are a little lower than NaClO₃'s melting temperature. When catalyzed by cobalt oxide, decomposition of the mixtures started at almost the
15 same temperatures as for NaClO₃ alone. Therefore, the alkaline additives can raise the decomposition temperature and provide smooth oxygen evolution.

Lithium aluminate, metaborate, tetraborate, phosphates, silicates and carbonate, calcium phosphates
20 and pyrophosphates strontium carbonate are all stable compounds and have very low solubility in water and they do not hydrate. This is a great advantage over other alkaline compounds. Therefore, oxygen generating formulations with these compounds can be either dry or wet
25 mixed. Oxygen generated by candles containing these compounds has a low moisture level.

Calcium phosphate and pyrophosphate are thermally stable. They are insoluble in water and do not hydrate. Both calcium phosphate and pyrophosphate are
30 non-toxic and have long been used in animal food to provide calcium. Therefore, these two calcium compounds are excellent additives to be used in sodium chlorate candles to smooth oxygen evolution and suppress chlorine. CaHPO₄ and Ca(H₂PO₄)₂ decompose at around 200°C to form
35 Ca₃(PO₄)₂ or Ca₂P₂O₇ and can thus be used.

Strontium carbonate is very stable in air. It does not decompose until 1170°C. It does not dissolve in water or hydrate. SrCO_3 inhibits to a less extent and raises the decomposition temperature just above the melting point of NaClO_3 . Therefore, candle containing SrCO_3 can have less fuel and the operation temperature will be lower, which is advantageous.

Lithium carbonate is chemically stable. It has low solubility in water and does not hydrate. At elevated temperatures it gradually decomposes to Li_2O and CO_2 . The decomposition is very slow in air below 850°C but much faster in molten NaClO_3 . However, lithium carbonate is a very effective inhibitor. As little as 0.2% Li_2CO_3 is enough to produce smooth oxygen evolution. Complete decomposition of the 0.2% Li_2CO_3 would produce only less than 2000 ppm CO_2 in the oxygen generated compared to allowed CO_2 concentration of 5000 ppm allowed in many applications.

Sodium metaborate and tetraborate, sodium metasilicate and orthosilicate are inexpensive and thermally very stable. Therefore, they are also good candidates as additives in the oxygen generator to smooth the decomposition and suppress chlorine formation.

The preferred candle composition includes about 0.05% to about 5.0% by weight of LiAlO_2 , LiBO_2 , Li_2CO_3 , Li_3PO_4 , Li_2SiO_3 , $\text{Li}_2\text{B}_4\text{O}_7$, Na_2SiO_3 , with 0.1% to 2.0% being particularly preferred and 0.05% to 10.0% $\text{Ca}_2\text{P}_2\text{O}_7$ or SrCO_3 , with about 0.2% to 4.0% by weight being particularly preferred. The chemicals purchased from Alfa Products or Aldrich Chemical Inc. have been found to be suitable. The same additives obtained from other sources are suitable so long as they have small particle size. If the additive chemicals as purchased are of relatively large particle size, these additives can also be made useable by grinding into fine powders.

The preferred catalysts include the cobalt oxides, such as CoO , Co_3O_4 , Co_2O_3 , or CoO_x , where x equals 1.0 to 1.5, and mixtures thereof. Cobalt oxide is particularly preferred as it can lower the decomposition temperature of sodium chlorate from 450°C to approximately 230°C . It may be obtained by decomposing cobalt carbonate or cobalt hydroxide. For example, cobalt carbonate may be decomposed at 260°C for 24 hours to produce a cobalt oxide with a specific surface area of approximately $100 \text{ m}^2/\text{g}$.

Preferred metal powder fuels include iron, manganese, cobalt, nickel, tungsten, titanium, magnesium, aluminum, niobium, zirconium, and mixtures thereof.

Iron, cobalt, or manganese powder having high purity and high surface area is especially preferred. Iron is particularly suitable. It may be obtained electrolytically or by hydrogen reduction. These processes yield iron powder substantially free of carbon and other impurities which would produce toxic compounds upon operation of the candle. Use of iron powder in the oxygen-generating candle furnishes heat and assists in stabilizing chlorate decomposition. Candles containing iron are less sensitive to surrounding temperatures than catalyzed fuel free candles previously described.

Preferred binders include glass powder, glass fiber, ceramic fiber, steel wool, and mixtures thereof. These binder materials can help to keep integrity of working candles and also help in mixing and pressing.

While the precise chemical reaction is not known, lithium carbonate, metaborate, tetraborate, phosphate and silicate, sodium metasilicate and strontium carbonate are basic and suppress free chlorine formation by either directly reacting chlorine gas or increasing the alkalinity of the candle and keeping chlorine gas from being formed.

The ignition pellet 52 typically may have a composition as follows:

	Fe	30%
	Fe ₂ O ₃	10%
5	Co ₂ O ₃	10%
	Glass powder	5%
	NaClO ₃	45%

10

EXAMPLE 1

1st Layer: 10.6 gram. 79.0% NaClO₃, 10.0% Fe, 7.0% CoO_x and 4.0% glass powder

15

2nd Layer: 34.3 gram. 86.9% NaClO₃, 7.5% Fe, 0.5% Li₂B₄O₇, 2.1% CoO_x and 3.0% glass powder

3rd Layer: 60.9 gram. 89.7% NaClO₃, 6.0% Fe, 0.5% Li₂B₄O₇, 0.8% CoO_x and 3.0% glass powder

20

4th Layer: 120.2 gram. 93.4% NaClO₃, 3.0% Fe, 0.5% Li₂B₄O₇, 0.6% CoO_x, and 2.5% glass powder

The candle lasted for 16.2 minutes with a oxygen release of 68.5 liters at room temperature by weight change. The impurities levels are 1,080 ppm CO₂, 1 ppm CO, less than 20 ppb chlorine, and 6.6 mg water per liter oxygen. The oxygen flow over time is shown in Figure 3.

30

EXAMPLE 2

1st Layer: 10.6 gram. 79.0% NaClO₃, 10.0% Fe, 9.0% CoO_x and 2.0% glass powder

35

2nd Layer: 34.3 gram. 86.5% NaClO₃, 8.0% Fe, 0.5% LiBO₂, 3.0% CoO_x and 2.0% glass powder

3rd Layer: 178.4 gram. 89.5% NaClO₃, 6.0% Fe, 0.5% LiBO₂, 2.0% CoO_x and 2.0% glass powder

40

The candle lasted for 10.5 minutes with a oxygen release of 64.9 liters at room temperature by weight change. The impurity levels are 995 ppm CO₂, 1 ppm CO, less than 20 ppb chlorine, and 6.4 mg water per liter

oxygen. The oxygen flow rate over time is shown in Figure 4.

5

EXAMPLE 3

- 1st Layer: 10.6 gram. 80.0% NaClO_3 , 10.0% Fe, 7.0% CoO_x and 3.0% glass powder
- 10 2nd Layer: 34.3 gram. 87.9% NaClO_3 , 7.5% Fe, 1.0% Li_3PO_4 , 1.6% CoO_x and 2.0% glass powder
- 3rd Layer: 60.9 gram. 90.2% NaClO_3 , 6.0% Fe, 1.0% Li_3PO_4 , 0.8% CoO_x and 2.0% glass powder
- 15 4th Layer: 120.2 gram. 93.6% NaClO_3 , 3.0% Fe, 1.0% Li_3PO_4 , 0.4% CoO_x , and 2.0% glass powder

20

The candle lasted for 18.2 minutes with a oxygen release of 67.0 liters at room temperature by weight change. The oxygen contains 1 ppm CO, 920 ppm CO_2 , 0 ppb Cl_2 , and 4.8 mg H_2O per liter oxygen. The oxygen flow rate over time is shown in Figure 5.

25

EXAMPLE 4

- 1st Layer: 10.6 gram. 77.0% NaClO_3 , 11.0% Fe, 9.0% CoO_x and 3.0% glass powder.
- 30 2nd Layer: 34.3 gram. 87.7% NaClO_3 , 7.5% Fe, 0.2% Li_2CO_3 , 1.6% CoO_x and 3.0% glass powder
- 3rd Layer: 60.9 gram. 90.0% NaClO_3 , 6.0% Fe, 0.2% Li_2CO_3 , 0.8% CoO_x and 3.0% glass powder.
- 35 4th Layer: 80.8 gram. 94.4% NaClO_3 , 3.0% Fe, 0.2% Li_2CO_3 , 0.4% CoO_x and 2.0% glass powder
- 40 5th Layer: 41.4 gram. 96.2% NaClO_3 , 1.3% Fe, 0.2% Li_2CO_3 , 0.3% CoO_x and 2.0% glass powder

The candle lasted for 15.3 minutes with a oxygen release of 69.3 liters at room temperature by weight

change. The impurity levels are 1490 ppm CO₂, 1 ppm CO, less than 20 ppb chlorine, 5.0 mg water per liter oxygen, and the oxygen evolution was smooth.

5

EXAMPLE 5

1st Layer: 10.6 gram. 81.0% NaClO₃, 9.0% Fe, 8.0% CoO_x and 2.0% glass powder

10

2nd Layer: 34.3 gram. 85.5% NaClO₃, 8.0% Fe, 0.5% Na₂SiO₃, 4.0% CoO_x and 2.0% glass powder

3rd Layer: 60.9 gram. 90.9% NaClO₃, 5.5% Fe, 0.5% Na₂SiO₃, 2.1% CoO_x and 2.0% glass powder

15

4th Layer: 120.3 gram. 92.7% NaClO₃, 3.5% Fe, 0.5% Na₂SiO₃, 1.5% CoO_x and 2.0% glass powder

20

The candle lasted for 14.7 minutes with a oxygen release of 67.8 liters at room temperature by weight change. The oxygen flow over time is given in Figure 6. The impurity levels of the oxygen are 0 ppm CO, 1530 ppm CO₂, 0 ppb Cl₂, and 7.3 mg H₂O per liter oxygen.

25

EXAMPLE 6

1st Layer: 10.6 gram. 77.0% NaClO₃, 11.0% Fe, 9.0% CoO_x and 3.0% glass powder

30

2nd Layer: 34.3 gram. 87.8% NaClO₃, 8.0% Fe, 0.5% Li₂CO₃, 1.7% CoO_x and 2.0% glass powder

3rd Layer: 60.9 gram. 90.7% NaClO₃, 6.0% Fe, 0.5% Li₂CO₃, 0.8% CoO_x and 2.0% glass powder

35

4th Layer: 83.7 gram. 94% NaClO₃, 3.1% Fe, 0.5% Li₂CO₃, 0.4% CoO_x and 2.0% glass powder

5th Layer: 36.4 gram. 96.0% NaClO₃, 1.2% Fe, 0.5% Li₂CO₃, 0.3% CoO_x and 2.0% glass powder

40

The candle lasted for 21.2 minutes with a oxygen

release of 69.2 liters at room temperature by weight change. The oxygen flow over time is shown in Figure 7.

EXAMPLE 7

- 5 1st Layer: 10.6 gram. 80.0% NaClO_3 , 10.0% Fe, 7.0% CoO_x
and 3.0% glass powder
- 2nd Layer: 34.3 gram. 87.3% NaClO_3 , 7.5% Fe, 0.5%
10 Li_2SiO_3 , 1.7% CoO_x and 3.0% glass powder
- 3rd Layer: 60.9 gram. 89.5% NaClO_3 , 6.0% Fe, 0.5%
15 Li_2SiO_3 , 1.0% CoO_x and 3.0% glass powder
- 4th Layer: 120.3 gram. 92.7% NaClO_3 , 3.5% Fe, 0.5%
 Li_2SiO_3 , 0.8% CoO_x and 2.5% glass powder

The candle lasted for 14.5 minutes with a oxygen
release of 66.2 liters at room temperature by weight
change and the oxygen evolution was smooth. The impurity
20 levels are 7 ppm CO, 580 ppm CO_2 , 0 ppb Cl_2 , and 6.1 mg H_2O
per liter oxygen.

EXAMPLE 8

- 1st Layer: 10.6 gram. 80.0% NaClO_3 , 10.0% Fe, 7.0% CoO_x
and 3.0% glass powder
- 25 2nd Layer: 34.3 gram. 85.9% NaClO_3 , 7.5% Fe, 2.0%
 $\text{Ca}_2\text{P}_2\text{O}_7$, 1.6% CoO_x and 3.0% glass powder
- 3rd Layer: 60.9 gram. 88.5% NaClO_3 , 6.0% Fe, 2.0%
30 $\text{Ca}_2\text{P}_2\text{O}_7$, 0.5% CoO_x and 3.0% glass powder
- 4th Layer: 120.3 gram. 92.5% NaClO_3 , 3.2% Fe, 2.0%
 $\text{Ca}_2\text{P}_2\text{O}_7$, 0.3% CoO_x and 2.0% glass powder

35 The candle lasted for 12.4 minutes with a oxygen
release of 67.4 liters at room temperature by weight
change and the oxygen evolution was smooth. The impurity
levels are 8 ppm CO, 950 ppm CO_2 , 0 ppb Cl_2 , and 6.4 mg H_2O
per liter oxygen.

Claims:

1. In a oxygen-generating candle composition that includes an amount of an alkali metal chlorate or perchlorate oxygen source other than lithium perchlorate alone and which upon initiation and decomposition thereof will yield oxygen and residual chlorine, the improvement which comprises a non-toxic additive incorporated into the candle composition for suppression of the residual chlorine and to enhance uniform oxygen generation and evolution, said additive being an alkaline compound selected from the group consisting of (a) a compound of the formula $M_aH_bX_cO_d$ where M = an alkali metal or alkaline earth metal, a = a number which is one or more, H = hydrogen, b = a number which is 0, or one or more, X = an element chosen from the group consisting of the elements of Group IIIA, the elements of Group IVA, the elements identified by the numbers 15 and 51 of Group VA, the elements of Group IVB, the elements of Group VB, the elements of Group VIB, and the element identified by the number 25 of Group VIIB of the Periodic Table, c = a number which is one or more, O = oxygen, and d = a number which is one or more; (b) lithium, sodium, potassium, rubidium, cesium, magnesium and strontium hydroxides; (c) magnesium, calcium and strontium peroxides; and (d) mixtures of the foregoing.

2. The oxygen-generating candle composition of Claim 1, wherein said additive is an alkaline compound selected from the group consisting of alkali metal and alkaline earth metal aluminates, borates, carbonates, chromates, germinates, hafnates, manganates, molybdenates, niobates, phosphates, silicates, stanates, tantalates, titanates, tungstates, vanadates, zirconates, and derivatives of the foregoing compounds in which the alkali metal and alkaline earth metal cations are partially substituted by hydrogen.

3. The oxygen-generating candle composition of Claim 1, wherein said additive is present in an amount of from about 0.05% to about 10% weight of the candle composition.

4. The oxygen-generating candle composition of Claim 3, wherein said additive is lithium tetraborate.

5. The oxygen-generating candle composition of Claim 3, wherein said additive is lithium metaborate.

6. The oxygen-generating candle composition of Claim 3, wherein said additive is lithium phosphate.

7. The oxygen-generating candle composition of Claim 3, wherein said additive is lithium carbonate.

8. The oxygen-generating candle composition of Claim 3, wherein said additive is lithium silicate.

5 9. The oxygen-generating candle composition of Claim 3, wherein the additive is a lithium compound provided in an amount of from about 0.05% to about 5% by weight of the candle composition.

10 10. The oxygen-generating candle composition of Claim 3, wherein the additive is a lithium compound provided in an amount of from about 0.1% to about 2% by weight of the candle composition.

15 11. The oxygen-generating candle composition of Claim 3, wherein the additive is a sodium compound provided in an amount of from about 0.05% to about 5% by weight of the candle composition.

20 12. The oxygen-generating candle composition of Claim 3, wherein the additive is a sodium compound provided in an amount of from about 0.1% to about 2% by weight of the candle composition.

25 13. The oxygen-generating candle composition of Claim 3, wherein the additive is a calcium compound provided in an amount of from about 0.05% to about 10% by weight of the candle composition.

14. The oxygen-generating candle composition of Claim 3, wherein the additive is a calcium compound provided in an amount of from about 0.2% to about 4% by weight of the candle composition.

5

15. The oxygen-generating candle composition of Claim 3, wherein the additive is a strontium compound provided in an amount of from about 0.05% to about 10% by weight of the candle composition.

10

16. The oxygen-generating candle composition of Claim 3, wherein the additive is a strontium compound provided in an amount of from about 0.2% to about 4% by weight of the candle composition.

15

17. The oxygen-generating candle composition of Claim 1, wherein is further included a cobalt catalyst in the candle composition.

20

18. The oxygen-generating candle composition of Claim 17, wherein said cobalt catalyst is a cobalt oxide.

25

19. The oxygen-generating candle composition of Claim 18, wherein said cobalt catalyst is selected from the group consisting of CoO , Co_3O_4 , Co_2O_3 and Co_x , wherein x ranges from about 1 to about 1.5, and mixtures of the foregoing.

20. The oxygen-generating candle composition of Claim 19, wherein said cobalt catalyst is present in an amount of from about 0.1% to about 5% by weight of the candle composition.

5

21. The oxygen-generating candle composition of Claim 17, wherein is further included a metal fuel in the candle composition.

10

22. The oxygen-generating candle composition of Claim 21, wherein said metal fuel is a powder selected from the group consisting of iron, manganese, cobalt, nickel, tungsten, titanium, magnesium, aluminum, niobium, zirconium, and mixtures of the foregoing.

15

23. The oxygen-generating candle composition of Claim 22, wherein said fuel is powdered iron.

20

24. The oxygen-generating candle composition of Claim 21, wherein is further included a binder selected from the group consisting of glass powder, glass fibers, ceramic fibers, steel wool, and mixtures of the foregoing.

25. In a method of generating breathable oxygen as a result of the initiation and operation of a candle composition that includes a cobalt oxide catalyzed alkali metal chlorate or perchlorate oxygen source other than lithium perchlorate alone, the improved step of adding a non-toxic additive to the oxygen source in order to increase the decomposition temperature of the alkali metal chlorate or perchlorate to a level above the melting temperature thereof that assures steady oxygen generation, said step including introducing into the chlorate or perchlorate before initiation thereof, an alkaline compound selected from the group consisting of (a) a compound of the formula $M_aH_bX_cO_d$, where M = an alkali metal or alkaline earth metal, a = a number which is one or more, H = hydrogen, b = a number which is one or more, X = an element chosen from the group consisting of the elements of Group IIIA, the elements of Group IVA, the elements identified by the numbers 15 and 51 of Group VA, the elements of Group IVB, the elements of Group VB, the elements of Group VIB, and the element identified by the number 25 of Group VIIB of the Periodic Table, c = a number which is one or more, O = oxygen, and d = a number which is 0, one or more; (b) lithium, sodium, potassium, rubidium, cesium, magnesium and strontium hydroxides; (c) magnesium, calcium and strontium peroxides; and (d) mixtures of the foregoing.

26. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes adding a compound
5 selected from the group consisting of alkali metal and alkaline earth metal aluminates, borates, carbonates, chromates, germinates, hafnates, manganates, molybdenates, niobates, phosphates, silicates, stanates, tantalates, titanates, tungstates, vanadates, zirconates, and
10 derivatives of the foregoing compounds in which the alkali metal and alkaline earth metal cations are partially substituted by hydrogen.

27. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an
15 alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes introducing from about 0.05% to about 10% weight of said additive to the oxygen source.

28. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an
20 alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes incorporating a lithium compound selected from the groups (a) and (b) set forth
25 which is provided in an amount from about 0.05% to about 5% by weight of the candle composition.

29. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes incorporating a sodium compound selected from the groups (a) and (b) set forth which is provided in an amount from about 0.05% to about 5% by weight of the candle composition.

30. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes incorporating a potassium compound selected from the groups (a) and (b) set forth which is provided in an amount from about 0.05% to about 5% by weight of the candle composition.

31. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes incorporating a calcium compound selected from the groups (a) and (c) set forth which is provided in an amount from about 0.05% to about 10% by weight of the candle composition.

32. A method of generating breathable oxygen as set forth in Claim 25, wherein the step of introducing an alkaline compound into the alkali metal chlorate or perchlorate oxygen source includes incorporating a
5 strontium compound selected from the groups (a) and (c) set forth which is provided in an amount from about 0.05% to about 10% by weight of the candle composition.

1/5

Fig. 1

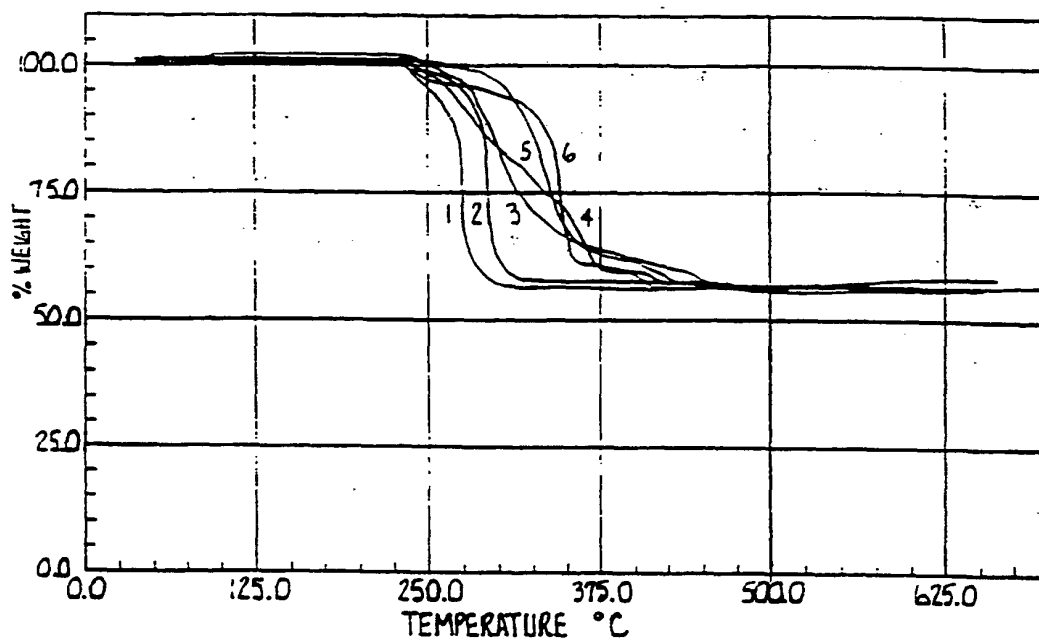
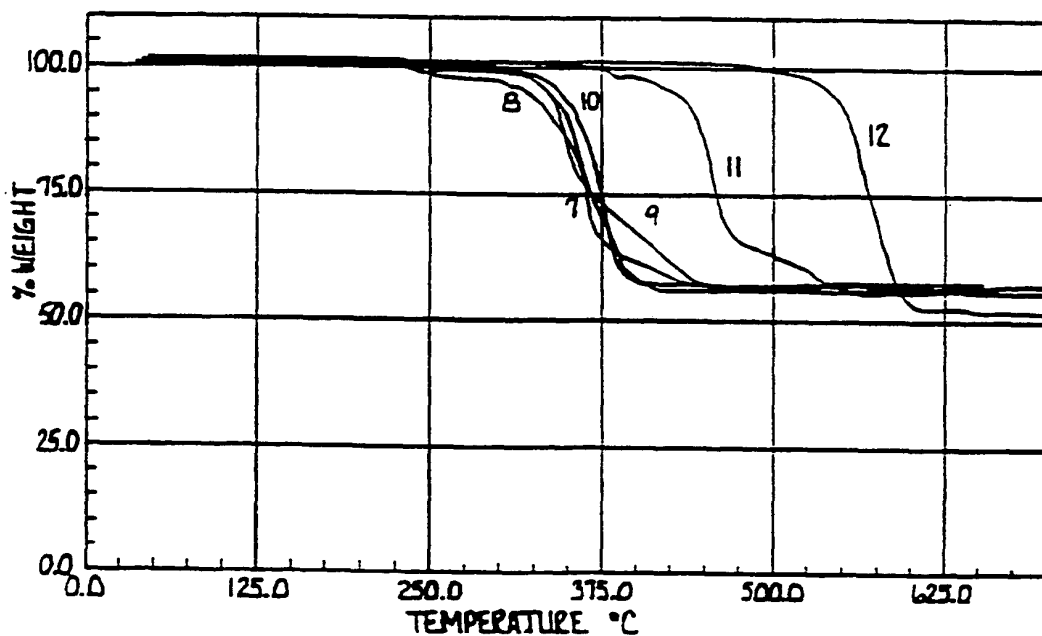
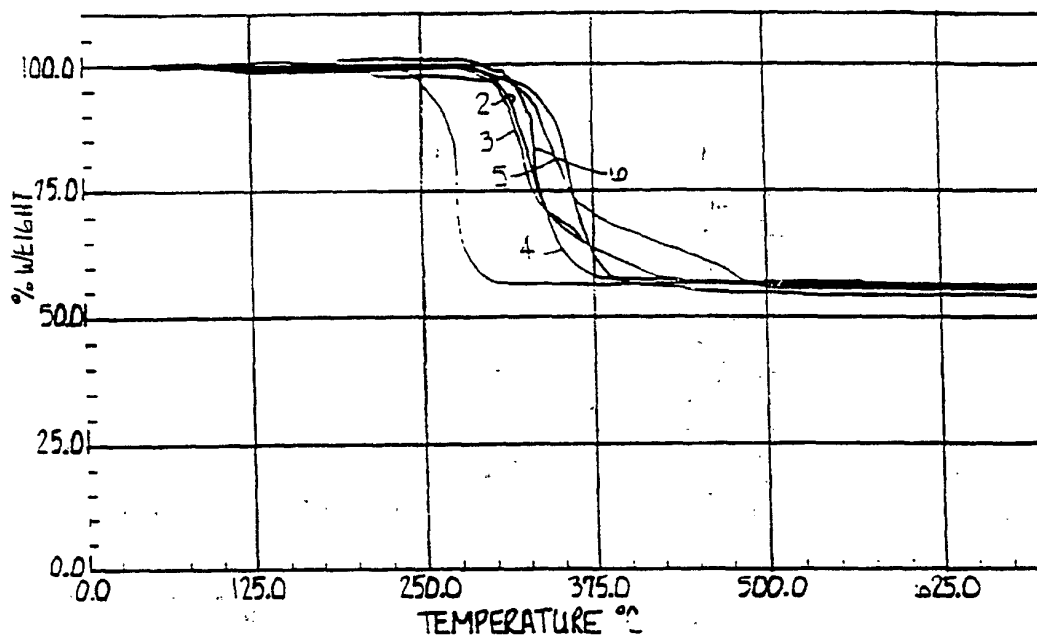
1) 99% NaClO₃ + 1% CoO_x3) 98% NaClO₃ + 1% Li₂B₄O₇ + 1% CoO_x5) 97% NaClO₃ + 2% Na₂SiO₃ + 1% CoO_x2) 95% NaClO₃ + 4% SrCO₃ + 1% CoO_x4) 97% NaClO₃ + 2% Ca₂P₂O₇ + 1% CoO_x6) 98% NaClO₃ + 1% Li₃PO₄ + 1% CoO_x

Fig. 1A

7) 98% NaClO₃ + 1% Ca(OH)₂ + 1% CoO_x9) 98.5% NaClO₃ + 0.5% Li₂O₂ + 1% CoO_x11) 98% NaClO₃ + 1% LiBO₂ + 1% CoO_x8) 98% NaClO₃ + 1% Li₂CO₃ + 1% CoO_x10) 95% NaClO₃ + 4% BaO₂ + 1% CoO_x12) 100% NaClO₃

2/5

Fig. 2

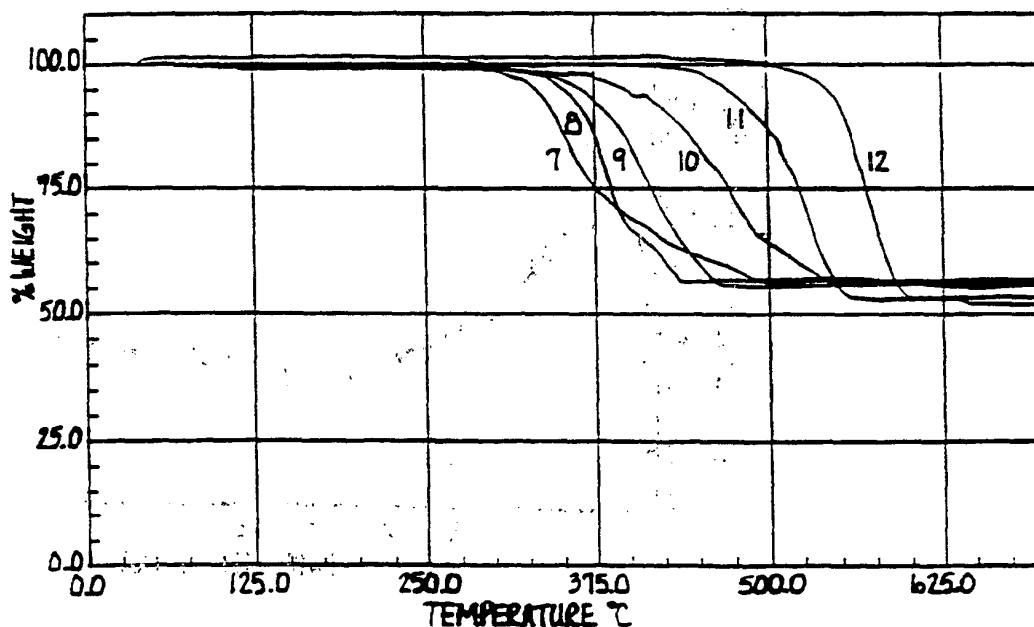


- 1.) 99% NaClO_3 + 1% CoO_x
 3.) 98% NaClO_3 + 1% NaClO_3 + 1% CoO_x
 5.) 96% NaClO_3 + 3% KOH + 1% CoO_x

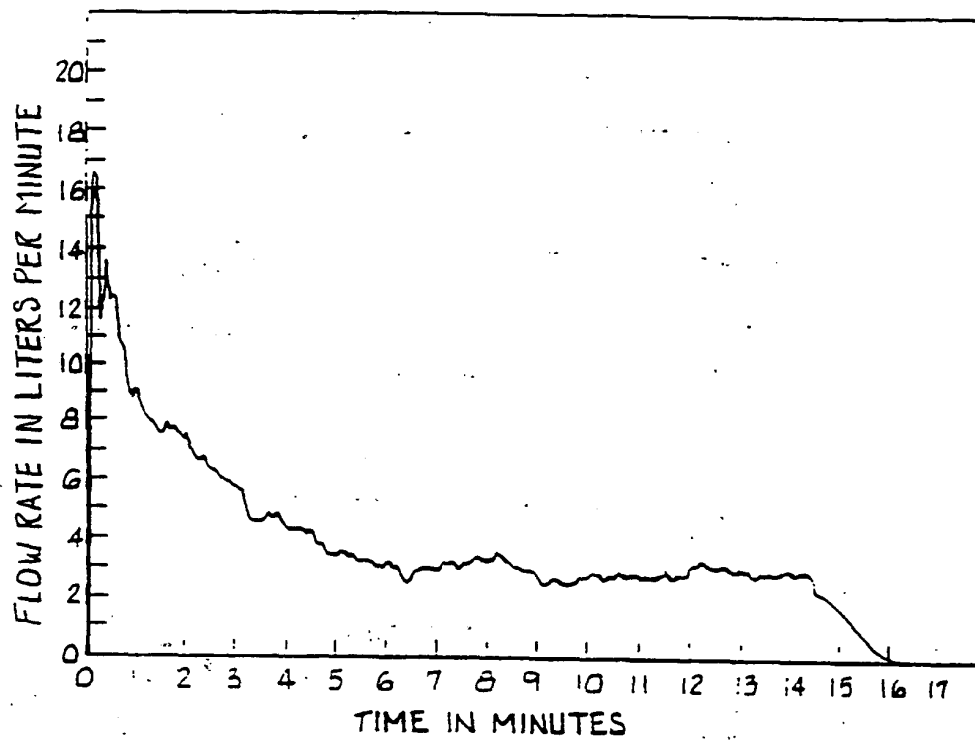
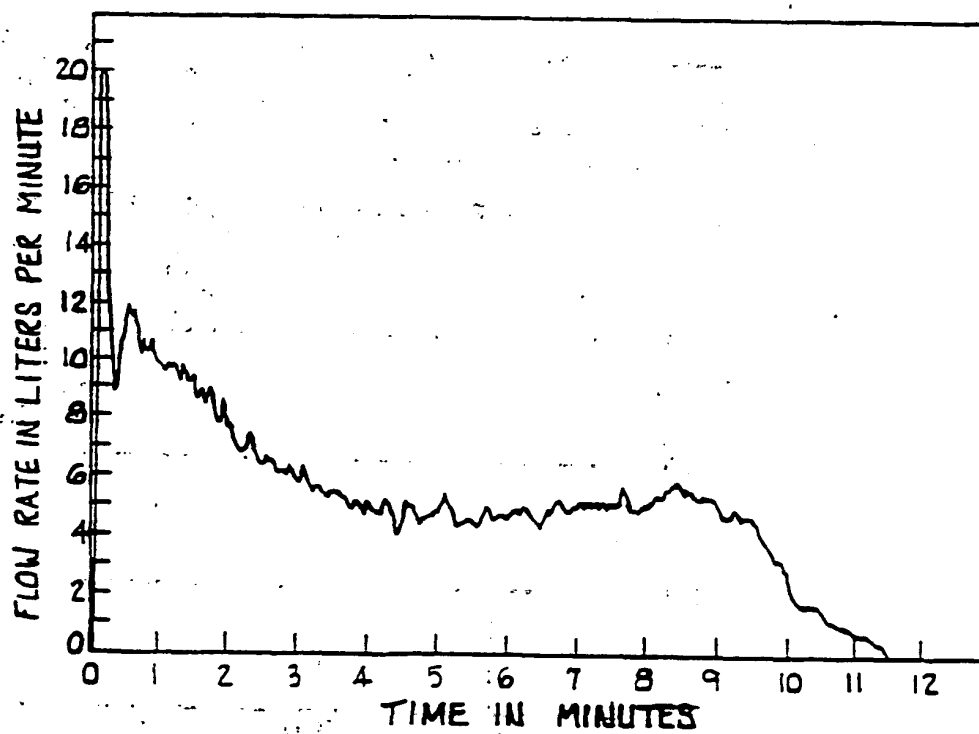
- 2.) 97% NaClO_3 + 2% NaHCO_3 + 1% CoO_x
 4.) 97% NaClO_3 + 2% $\text{Na}_2\text{Cr}_2\text{O}_7 \cdot 2\text{H}_2\text{O}$ + 1% CoO_x
 6.) 96% NaClO_3 + 3% $\text{Na}_3\text{PO}_4 \cdot 12\text{H}_2\text{O}$ + 1% CoO_x

Fig. 2 A

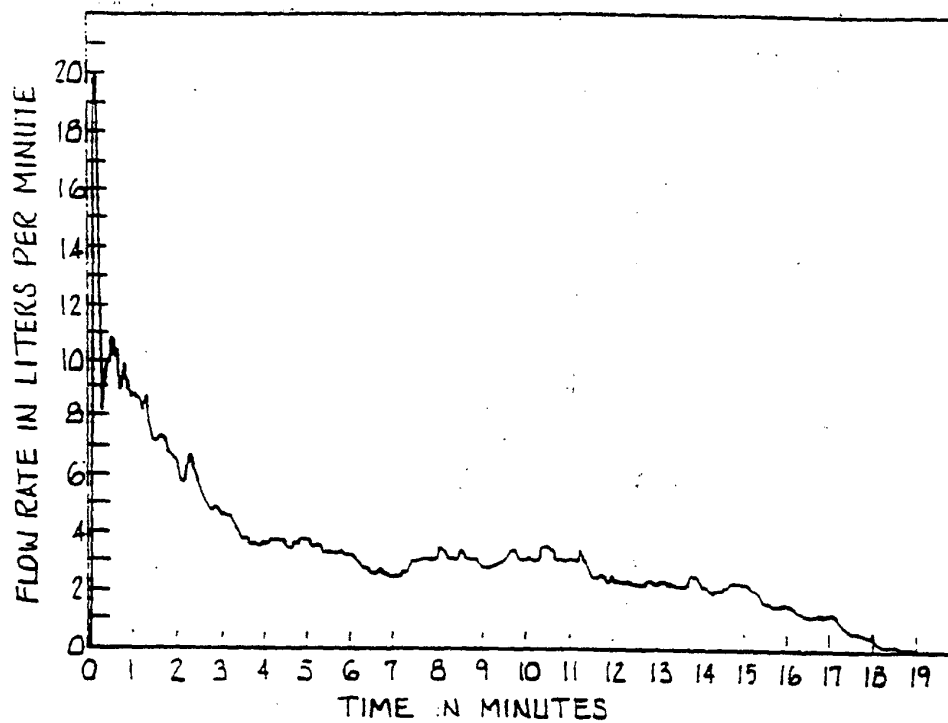
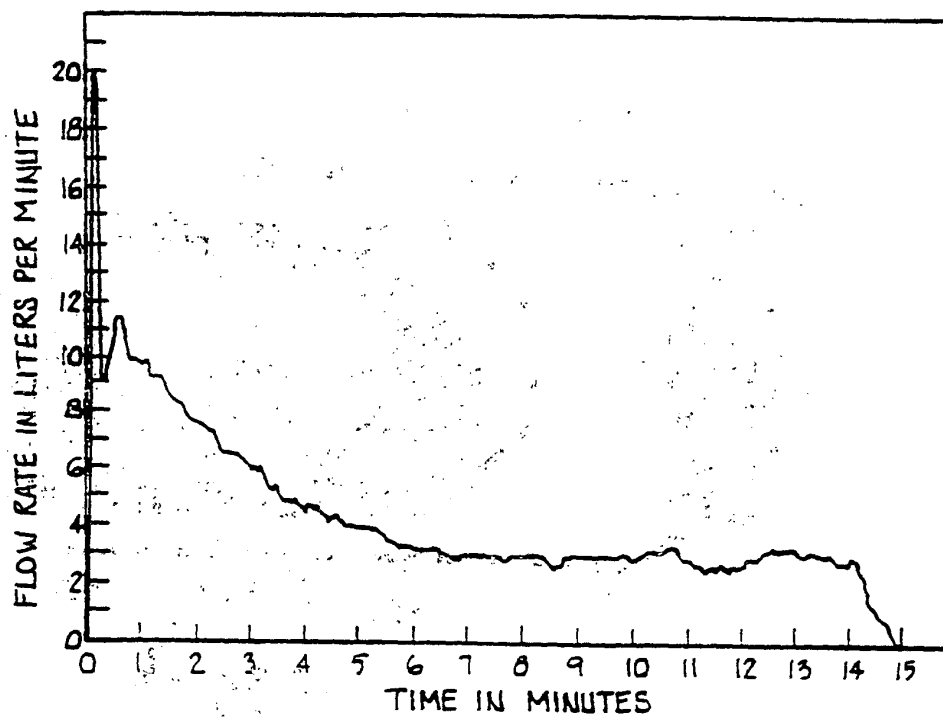
- 7.) 96% NaClO_3 + 3% NaOH + 1% CoO_x
 9.) 97% NaClO_3 + 2% LiOH + 1% CoO_x
 11.) 97% NaClO_3 + 2% $\text{Sr}(\text{OH})_2 \cdot \text{H}_2\text{O}$ + 1% CoO_x
 8.) 95% NaClO_3 + 4% $\text{Ba}(\text{OH})_2 \cdot \text{H}_2\text{O}$ + 1% CoO_x
 10.) 97% NaClO_3 + 2% Na_2BO_2 + 1% CoO_x
 12.) 100% NaClO_3



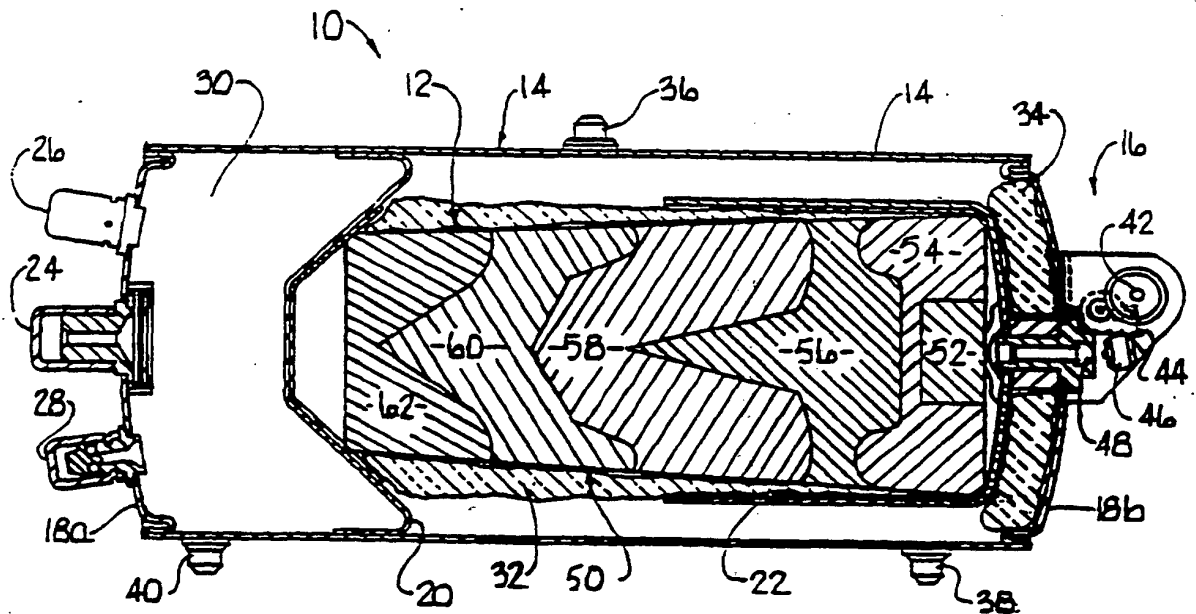
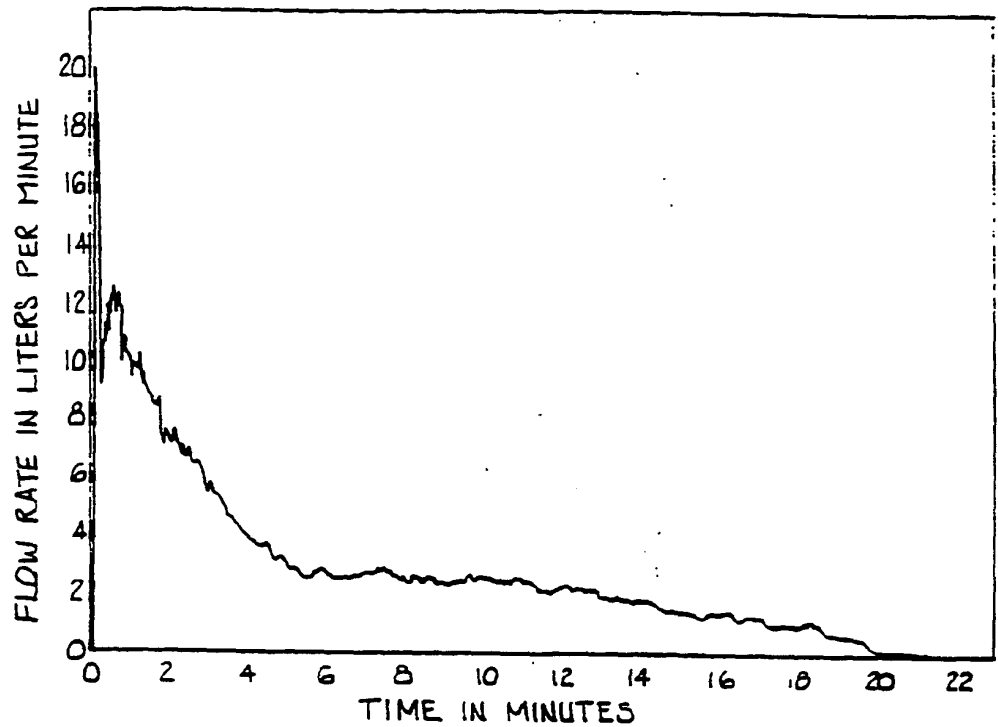
3/5

Fig. 3*Fig. 4*

4/5

Fig. 5*Fig. 6*

5/5

Fig. 7*Fig. 8*

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US93/01562

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) : C01B 11/14; C01B 11/18

US CL : 252/187.31

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 252/187.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US, A, 2,004,243 (HLOCH) 11 June 1935, See claims 1-7.	1,3,11-16, 21-25, & 27-30
A	US, A, 3,702,305 (THOMPSON) 07 November 1972.	1,3,11-16, 21-25,27- 30
Y	CN, A, 1035248A (HAICHUN ET AL.) 06 September 1989, See the examples.	1,3,9-10, 17-25,27-28
Y	US, A, 3,293,187 (MARKOWITZ) 20 December 1966, See column 2, lines 19-25.	1,3,9-10 17-25,27-28

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

* Special categories of cited documents:	*T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A document defining the general state of the art which is not considered to be part of particular relevance	*X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E earlier document published on or after the international filing date	*Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combinations being obvious to a person skilled in the art
*L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*& document member of the same patent family
*O document referring to an oral disclosure, use, exhibition or other means	
*P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

29 APRIL 1993

Date of mailing of the international search report

27 MAY 1993

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. NOT APPLICABLE

Authorized officer

JOSEPH D. ANTHONY

Telephone No. (703) 308-0446

FULWIDER PATTON LEE & UTECHT, LLP
Attn. Paul, James W.
Howard Hughes Center,
6060 Center Drive, 10th Floor
Los Angeles, California 90045
UNITED STATES OF AMERICA

Date: 17/04/2002

